## COPY OF PAPERS ORIGINALLY FILED

1645

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			First Named Inventor	Levi	.n	JA	1282
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Fee Transmittal Form  Fee Attached  Amendment / Reply (Second Preliminary)  After Final  Affidavits/declaration(s)  Extension of Time Request  Express Abandonment Request  Information Disclosure Statement  Certified Copy of Priority Document(s)  Response to Missing Parts/ Incomplete Application  Response to Missing Parts under 37 CFR 1.52 or 1.53		Continue   Continue	to Convert to a snal Application of Attorney, Revocation of Correspondence	At of At	Group ppeal Communication Appeals and Interfer ppeal Communication ppeal Notice, Brief, Reply Proprietary Information Status Letter Other Enclosure(s) (Interfer) Interfer)	erences ion to Group Brief) ion	
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m dividual name	Mary Kakefuda, Reg. No. 39,245 Syngenta Biotechnology, Inc.						
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Burden Hour Statement: This form is estimated to take 0.2 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, DC 20231.

PE JC135

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Melissa Hardy

Type or print name

October 22, 2001

Date

## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

RECEIVED

IN RE APPLICATION OF

LEVIN ET AL.

**APPLICATION NO.: 09/896,186** 

FILED: June 29, 2001

FOR: Methods of Controlling Gene Expression

and Gene Silencing

ART UNIT: 1645

**CONFIRMATION NO.: 9567** 

DOCKET: PB/5-31481A

JAN 2 8 2002

**TECH CENTER 1600/290** 

## SECOND PRELIMINARY AMENDMENT UNDER 37 C.F.R. §1.111

Box: Non-Fee Amendment Commissioner for Patents Washington, D.C. 20231

Sir:

Prior to examination of the above-identified patent application, please enter the following amendments and consider the remarks below.

## IN THE SPECIFICATION:

Please amend the specification to read as follows:

On page 52, Example 5:

The line 8Z-2 (see Example 4 above) is crossed with the line S11.13-34 (see Example 2 above) and the resultant F1 generation plants are allowed to self-fertilize to obtain the F2 generation. Approximately 60 F2 plants are grown and tested for a presence of T-DNA insertion in the RDRD gene derived from the S11.13-34 parental line and for the 35S-GFP T-DNA derived from the 8Z-2 parental line. The presence of the T-DNA insertion in the RDRD gene is demonstrated as described in Example 2. Plants carrying this T-DNA insertion are then checked for homozygosity by PCR using the 3' specific primer (SEQ ID NO:19) and 36851TD#3 (5'-gct ccg ccc aca taa ttc aaa caa cac-3', SEQ ID NO:29). These primers span a region of genomic DNA including the insertion site such that only the wild-type copy of DNA results in